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2. REPORT DATE 19 March 1993		3. REPORT TYPE AND DATES COVERED Technical Report	
4. TITLE AND SUBTITLE π - π Interaction in the Complexes of the Pyridino-18-crown-6 Host Molecules and Aromatic Ammonium Salts as Determined by ^1H NMR Spectroscopy		5. FUNDING NUMBERS N00014-91-J-1710 R & T Code 413p002	
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9. SPONSORING / MONITORING AGENCY NAME(S) AND ADDRESS(ES) Dr. H. Guard Office of Naval Research 800 North Quincy Street Arlington, VA 22217-5000		10. SPONSORING / MONITORING AGENCY REPORT NUMBER N/A	
11. SUPPLEMENTARY NOTES			
12a. DISTRIBUTION / AVAILABILITY STATEMENT Approved for public release; distribution unlimited		12b. DISTRIBUTION CODE	
13. ABSTRACT (Maximum 200 words) Significant π - π interaction is found in the complexes of (S,S)-dimethylpyridino-18-crown-6 with (R)- and (S)-[α -(1-Naphthyl)ethyl]ammonium perchlorate. This finding is supported by the ^1H NOESY NMR spectral technique, greater chemical shift changes of aromatic protons in both host and guest molecules upon complexation, and by molecular mechanics calculations.			
14. SUBJECT TERMS		15. NUMBER OF PAGES	
		16. PRICE CODE N/A	
17. SECURITY CLASSIFICATION OF REPORT Unclassified	18. SECURITY CLASSIFICATION OF THIS PAGE Unclassified	19. SECURITY CLASSIFICATION OF ABSTRACT Unclassified	20. LIMITATION OF ABSTRACT UL

OFFICE OF NAVAL RESEARCH

Grant N00014-91-J-1710

R&T Code 413p002

TECHNICAL REPORT NO. 18

π - π Interaction in the Complexes of the Pyridino-18-crown-6 Host
Molecules and Aromatic Ammonium Salts as Determined by ^1H NMR Spectroscopy

by

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March 19, 1993

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Enantiomeric recognition of chiral organic ammonium salts by many of these chiral macrocyclic ligands has been studied by NMR spectroscopy, calorimetric titration, solvent extraction, liquid membrane transport and chromatography. Detailed discussions of the synthesis and characterization of chiral ligands are given in a few reviews.¹⁻⁵ Among the many chiral crown ethers, chiral pyridino-18-crown-6 ligands display moderate to significant recognition.^{4,5} However, interactions of most of the reported chiral ligands with enantiomeric guests have not been well characterized from either a thermodynamic or a kinetic standpoint.⁵ Structural and conformational studies of complexes in solution could provide evidence for the basis of enantiomeric recognition.

The primary binding force between the pyridino-crown ligand and the primary ammonium cation is that formed by the three hydrogen bonds as shown in Figure 1.^{6,7} However, the π - π interaction between the naphthene group of NapEt and the pyridine ring of the chiral ligand is considered as one of the possible causes for enantiomeric recognition.^{8,9} It is important to know if a π - π interaction exists and if it plays a role in promoting enantiomeric recognition in these chiral complexes. We now report an analysis of the π - π interaction in the complexes of (*S,S*)-Me₂P18C6 with (*R*)- and (*S*)- NapEt in a 50%CD₃OD and 50%CDCl₃ (v/v) solvent system by 2D ¹H NOESY spectroscopy and chemical shift changes.

Experimental

NOESY Experiments: 2D ¹H NOESY spectra of (*S,S*)-Me₂P18C6 complexes with (*R*)- and (*S*)-NapEt were taken on a Varian VXR-500 MHz NMR spectrometer at 25 °C. Stoichiometric amounts of host and guest were weighed on a high precision balance and dissolved in a mixed

solvent of 50%CD₃OD and 50%CDCl₃ (v/v) to give a concentration of 0.040 M. The 90 degree pulse width was always calibrated. Delay time was set to about 3 times the maximum T1 value.

Results and Discussion

When forming host-guest complexes with various ammonium salts in different solvent systems, the pyridine protons of the host showed significant chemical shift changes. A similar phenomenon has been observed in other systems.⁹ Table 1 lists the chemical shift values for free and complexed (*S,S*)-Me₂P18C6 with (*R*)-NapEt [(*S,S*)-(*R*)] and (*S*)-NapEt [(*S,S*)-(*S*)] as well as the differences between them. From these data, we can see that the pyridine proton in position 1 (Figure 1) has a 0.19 ppm upfield shift and a 0.01 ppm downfield shift in the of complexes with (*R*)-NapEt and (*S*)-NapEt, respectively. Proton 2 has 0.35 ppm and 0.10 ppm upfield shifts for (*S,S*)-(*R*) and (*S,S*)-(*S*) complexes, respectively. Significant changes (0.33 and 0.12 ppm upfield shifts) also occurred at position 4 which is close to the pyridine ring. Chemical shift changes can be caused by many factors such as hydrogen bonding, conformation changes, shielding and deshielding effects of aromatic rings, and π - π interactions. Hydrogen bonding and conformation changes should have little effect on the pyridine protons because the protons on the pyridine ring are not involved at the complexation site and the ring is a geometrically stable moiety. Deshielding and shielding effects of the π - π interacting units are thus expected to play an important role in the chemical shift changes of the pyridine protons. The chemical shift changes shown in Table 1 suggest the presence of a strong π - π interaction between the pyridine and naphthalene rings. This conclusion is supported by the chemical shift changes of guest molecule NapEt (Table 2). The

chemical shift assignments were made from an ^1H COSY spectrum. In this case, significant chemical shift changes occurred in positions A, C, E and F in the (*S,S*)-(*R*) complex and B, C and G in the (*S,S*)-(*S*) complex (see NapEt, Figure 1, for proton assignments). The relatively large chemical shift changes indicate that these positions are affected by the pyridine ring. The presence of positive and negative chemical shift changes results from shielding and deshielding effects of the pyridine ring depending on the relationship to the face or edge of the ring. Unlike a fixed structure in the crystalline state,^{7,8} these data do not determine an absolute geometry for the interaction.

The data in Tables 1 and 2 also show that the protons in the (*S,S*)-(*R*) complex have a greater average chemical shift change than those in the (*S,S*)-(*S*) complex. Although the size of the absolute value of chemical shift change for a specific proton is not necessarily proportional to the stability of a complex,¹⁰ the greater overall changes in the (*S,S*)-(*R*) complex over the (*S,S*)-(*S*) complex indicate greater stability for the (*S,S*)-(*R*) complex.

The presence of a π - π interaction is also strongly supported by data from 2D ^1H NOESY spectra (Figure 2). In the NOESY spectra, there are strong off-diagonal signals in the 7.2-8.2 ppm range between the chemical shifts of the pyridine and naphthalene protons indicating that pyridine protons of the ligand and naphthalene protons of the ammonium salt are spatially close and that the complex is stable enough for NOE cross-relaxation to occur.¹¹ As shown in Figure 2, the π systems overlap spatially in both the (*S,S*)-(*R*) (Figure 2a) and (*S,S*)-(*S*) (Figure 2b) complexes. This π - π interaction helps bind the two aromatic systems together, determines the steric environment for the methyl groups, and contributes to recognition.

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Table 1. ^1H NMR Chemical Shift Values (ppm) for (*S,S*)-Me₂P18C6 and Its Complexes with (*R*) & (*S*)-NapEt in 1M/1C^a Mixed Solvent System (v/v) at 25°C

Proton ^b	(<i>S,S</i>)-Me ₂ P18C6	(<i>S,S</i>)-(<i>R</i>) Complex ^c		(<i>S,S</i>)-(<i>S</i>) Complex ^d	
	ppm	ppm	Δ ppm	ppm	Δ ppm
1	7.75	7.56	-0.19	7.76	+0.01
2	7.34	6.99	-0.35	7.24	-0.10
4	4.80	4.47	-0.33	4.68	-0.12
5	3.81	3.81	0.00	3.80	-0.01
6	3.47	3.54	+0.07	3.41	-0.06
7&8 ^e	3.59-3.48	3.78-3.68		3.62-3.48	
9	1.19	1.25	+0.06	1.18	-0.01

^a M = CD₃OD, C = CDCl₃

^b Numbers correspond to protons in (*S,S*)-Me₂P18C6 (see structure)

^c (*S,S*)-(*R*) = Complex of (*S,S*)-Me₂P18C6 with (*R*)-NapEt

^d (*S,S*)-(*S*) = Complex of (*S,S*)-Me₂P18C6 with (*S*)-NapEt

^e Estimated values because of overlapping peaks.

Table 2. ¹H NMR Chemical Shift Values (ppm) for (*R*) & (*S*) NapEt and Their Complexes with (*S,S*)-Me₂P18C6 in 1M/1C^a Mixed Solvent System (v/v) at 25°C

Proton ^b	NapEt	(<i>S,S</i>)-(<i>R</i>) Complex ^c		(<i>S,S</i>)-(<i>S</i>) Complex ^d	
	ppm	ppm	Δppm	ppm	Δppm
A	7.63	7.78	+0.15	7.66	+0.03
B	7.65	7.61	-0.04	7.49	-0.16
C	8.07	7.94	-0.13	7.91	-0.16
D	7.92	7.84	-0.08	7.94	+0.02
E or F	7.59	7.25	-0.34	7.57	-0.02
F or E	7.59	7.40	-0.19	7.57	-0.02
G	7.94	7.84	-0.10	8.06	+0.12
K	5.34	5.35	+0.01	5.30	-0.04
L	1.79	1.84	+0.05	1.83	+0.04

^a See footnote a in Table 1

^b Letters correspond to protons in NapEt (see structure)

^c See footnote c in Table 1

^d See footnote d in Table 1

Figure 2a

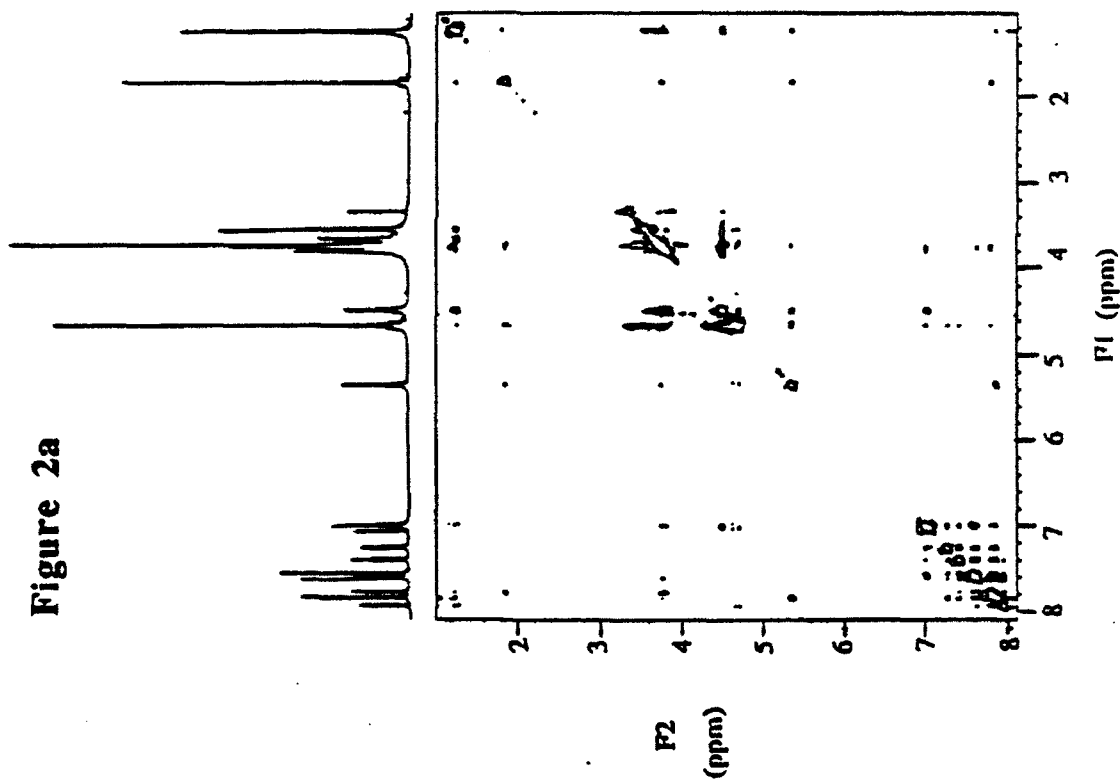


Figure 2b

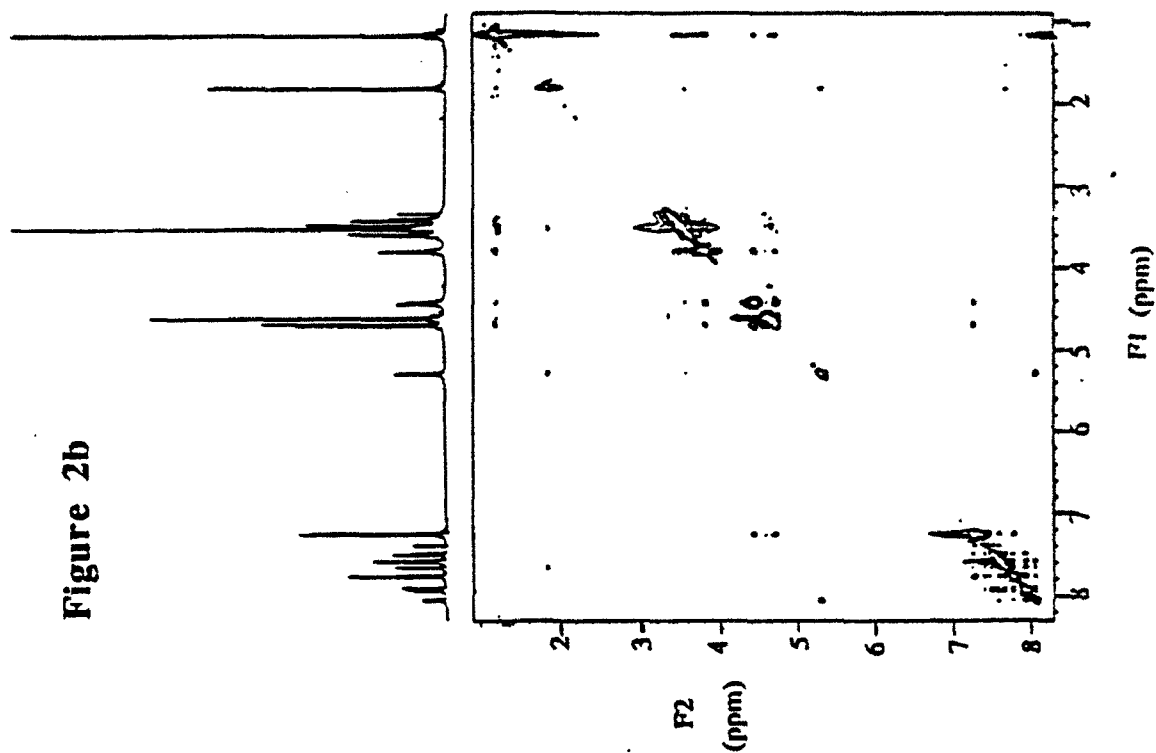
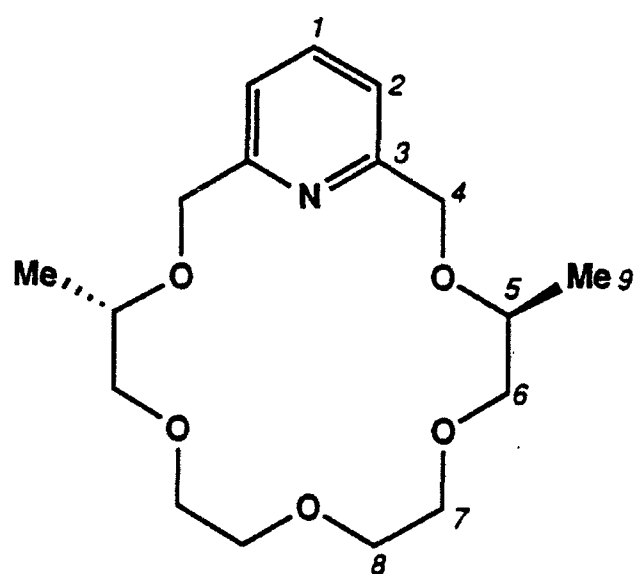
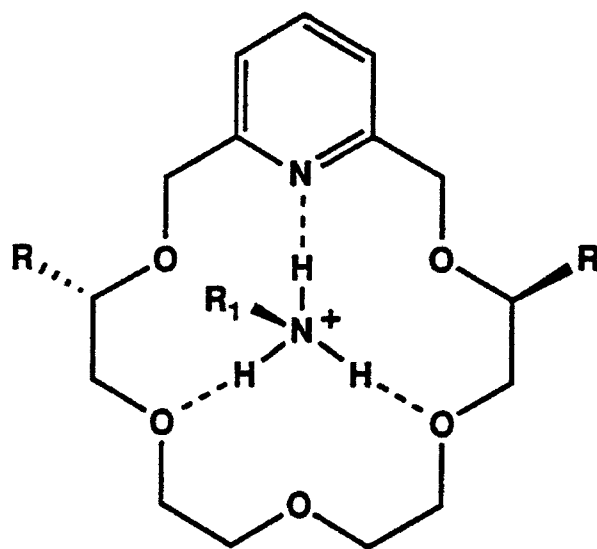
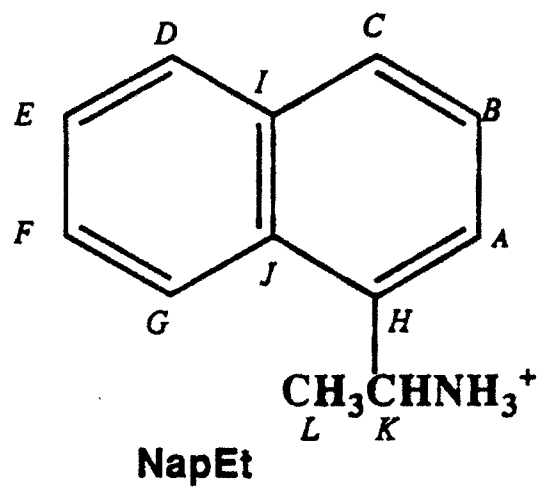


Figure 2. ^1H NMR NOESY spectra of complexes of (S,S) -dimethylpyridino-18-crown-6 with (R) -[α -(1-Naphthyl)ethyl]ammonium perchlorate (Figure 2a) and (S) -perchlorate (Figure 2b) in a mixed solvent of 50% CD_3OD + 50% CDCl_3 at 25°C

Figure 1. Structures of Compounds



Me₂P18C6



Tripoa